



## Inventor Name Search

Enter the **first few letters** of the Inventor's Last Name.  
Additionally, enter the **first few letters** of the Inventor's First name.

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Day : Thursday  
Date: 5/11/2006

Time: 14:04:15

## Inventor Name Search

Enter the **first few letters** of the Inventor's Last Name.  
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Last Name	First Name	
<input type="text" value="Wadgaonkar"/>	<input type="text" value="Dilip"/>	<input type="button" value="Search"/>

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## Refine Search

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### Search Results -

Terms	Documents
Dilip near Wadgaonkar	1

Database:

US Pre-Grant Publication Full-Text Database  
 US Patents Full-Text Database  
 US OCR Full-Text Database  
 EPO Abstracts Database  
 JPO Abstracts Database  
 Derwent World Patents Index  
 IBM Technical Disclosure Bulletins

Search:

L4






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### Search History

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 DATE: Thursday, May 11, 2006    [Printable Copy](#)    [Create Case](#)

Set Name Query

side by side

Hit Count Set Name

result set

*DB=PGPB,USPT; PLUR=YES; OP=OR*

L4	Dilip near Wadgaonkar	1	<a href="#">L4</a>
L3	Divya near Tewari	1	<a href="#">L3</a>
L2	David near Dixon	65	<a href="#">L2</a>
L1	Vijai near Kumar	21	<a href="#">L1</a>

END OF SEARCH HISTORY

(FILE 'HOME' ENTERED AT 14:13:55 ON 11 MAY 2006)

FILE 'CAPLUS, MEDLINE' ENTERED AT 14:14:02 ON 11 MAY 2006

L1 67998 S ANALGESIC  
L2 0 S L1 AND GEL(5A) (FORM?(3A) POLYMER)  
L3 130 S L1 AND ((POLYVINYL(1A)ALCOHOL) OR (POLYETHYLENE(1A)OXIDE) OR  
L4 0 S L3 AND EMETIC  
L5 149 S L1 AND EMETIC  
L6 0 S L5 AND (NASAL (5A) IRRITANT)  
L7 1 S L5 AND IRRITANT  
L8 1 S L3 AND (ZINC(2A) (SULFATE OR SULPHATE))  
L9 127 DUPLICATE REMOVE L3 (3 DUPLICATES REMOVED)  
L10 127 FOCUS L9 1-  
L11 10 S L10 AND (POLOXAMER OR (SODIUM(W)LAURYL(W)SULFATE) OR (SODIUM(  
L12 12040 S L1 AND OPIOID  
L13 28 S L12 AND EMETIC  
L14 24 DUPLICATE REMOVE L13 (4 DUPLICATES REMOVED)  
L15 24 FOCUS L14 1-  
L16 0 S L12 AND (ZINC(W) (SULFATE OR SULPHATE))

L7 ANSWER 1 OF 1 CAPLUS COPYRIGHT 2006 ACS on STN  
TI Pharmaceutical composition with an opiate and an **irritant**  
AB An oral composition comprising about 0.1-40% of an opiate and about 0.001-85% of an **irritant**, e.g., a capsaicinoid, an **emetic**, or histamine, for treating pain is described. A method is also described for discouraging abuse of an opiate comprising combining a therapeutically effective amount of an opiate and an **irritant** into an oral dosage form.

ACCESSION NUMBER: 2003:376651 CAPLUS  
DOCUMENT NUMBER: 138:390921  
TITLE: Pharmaceutical composition with an opiate and an **irritant**  
INVENTOR(S): Anderson, Kirsten A.; Hoch, James M.; Duvall, Jean-Marie; Liversidge, Gary; Slavitt, Joshua R.  
PATENT ASSIGNEE(S): Elan Corporation, PLC, Ire.  
SOURCE: PCT Int. Appl., 20 pp.  
CODEN: PIXXD2  
DOCUMENT TYPE: Patent  
LANGUAGE: English  
FAMILY ACC. NUM. COUNT: 1  
PATENT INFORMATION:

PATENT NO.	KIND	DATE	APPLICATION NO.	DATE
WO 2003039561	A1	20030515	WO 2002-US35397	20021104
W: AE, AG, AL, AM, AT, AU, AZ, BA, BB, BG, BR, BY, BZ, CA, CH, CN, CO, CR, CU, CZ, DE, DK, DM, DZ, EC, EE, ES, FI, GB, GD, GE, GH, GM, HR, HU, ID, IL, IN, IS, JP, KE, KG, KP, KR, KZ, LC, LK, LR, LS, LT, LU, LV, MA, MD, MG, MK, MN, MW, MX, MZ, NO, NZ, OM, PH, PL, PT, RO, RU, SD, SE, SG, SI, SK, SL, TJ, TM, TN, TR, TT, TZ, UA, UG, US, UZ, VC, VN, YU, ZA, ZM, ZW, AM, AZ, BY, KG, KZ, MD, RU, TJ, TM				
RW: GH, GM, KE, LS, MW, MZ, SD, SL, SZ, TZ, UG, ZM, ZW, AT, BE, BG, CH, CY, CZ, DE, DK, EE, ES, FI, FR, GB, GR, IE, IT, LU, MC, NL, PT, SE, SK, TR, BF, BJ, CF, CG, CI, CM, GA, GN, GQ, GW, ML, MR, NE, SN, TD, TG				
CA 2464528	AA	20030515	CA 2002-2464528	20021104
US 2003125347	A1	20030703	US 2002-288262	20021104
EP 1450824	A1	20040901	EP 2002-789428	20021104
R: AT, BE, CH, DE, DK, ES, FR, GB, GR, IT, LI, LU, NL, SE, MC, PT, IE, SI, LT, LV, FI, RO, MK, CY, AL, TR, BG, CZ, EE, SK				
JP 2005508372	T2	20050331	JP 2003-541852	20021104
PRIORITY APPLN. INFO.:			US 2001-340111P	P 20011102
			WO 2002-US35397	W 20021104
REFERENCE COUNT:	3	THERE ARE 3 CITED REFERENCES AVAILABLE FOR THIS RECORD. ALL CITATIONS AVAILABLE IN THE RE FORMAT		

L11 ANSWER 5 OF 10 CAPLUS COPYRIGHT 2006 ACS on STN

TI Sustained-release opioid formulations for treating pain

AB The invention combines two different subunits with different release profiles in novel sustained-release oral dosage forms. In particular, the oral dosage forms with specified drug release rate include a subunit that comprises an opioid **analgesic** and a sustained-release material, providing a duration of therapeutic effect of about 24 h. For example, a sustained-release oral formulation comprised (i) a first subunit releasing opioid in a sustained manner beginning in the first 12 h after administration containing morphine sulfate 50.0 mg, nonpareil seed (#16-18 mesh) 131.9 mg, hypromellose 3.3 mg, Et cellulose 28.1 mg, polyethylene glycol 6000 9.9 mg, Eudragit L100-55 8.3 mg, di-Et phthalate 5.7 mg, and talc 26.0 mg, and (ii) a second subunit releasing opioid in a sustained manner beginning in the second 12 h after administration containing morphine sulfate 50.0 mg, nonpareil seed (#16-18 mesh) 131.9 mg, hypromellose 3.3 mg, Eudragit RS PO 26.9 mg, Eudragit RL PO 4.5 mg, tri-Et citrate 3.1 mg, **sodium lauryl sulfate** 0.7 mg, and talc 17.6 mg.

ACCESSION NUMBER: 2004:270017 CAPLUS

DOCUMENT NUMBER: 140:292653

TITLE: Sustained-release opioid formulations for treating pain

INVENTOR(S): Boehm, Garth; Liang, Alfred

PATENT ASSIGNEE(S): Alpharma, Inc., USA

SOURCE: PCT Int. Appl., 44 pp.

CODEN: PIXXD2

DOCUMENT TYPE: Patent

LANGUAGE: English

FAMILY ACC. NUM. COUNT: 1

PATENT INFORMATION:

PATENT NO.	KIND	DATE	APPLICATION NO.	DATE
WO 2004026256	A2	20040401	WO 2003-US29634	20030922
WO 2004026256	A3	20040701		
W:	AE, AG, AL, AM, AT, AU, AZ, BA, BB, BG, BR, BY, BZ, CA, CH, CN, CO, CR, CU, CZ, DE, DK, DM, DZ, EC, EE, EG, ES, FI, GB, GD, GE, GH, GM, HR, HU, ID, IL, IN, IS, JP, KE, KG, KP, KR, KZ, LC, LK, LR, LS, LT, LU, LV, MA, MD, MG, MK, MN, MW, MX, MZ, NI, NO, NZ, OM, PG, PH, PL, PT, RO, RU, SC, SD, SE, SG, SK, SL, SY, TJ, TM, TN, TR, TT, TZ, UA, UG, US, UZ, VC, VN, YU, ZA, ZM, ZW			
RW:	GH, GM, KE, LS, MW, MZ, SD, SL, SZ, TZ, UG, ZM, ZW, AM, AZ, BY, KG, KZ, MD, RU, TJ, TM, AT, BE, BG, CH, CY, CZ, DE, DK, EE, ES, FI, FR, GB, GR, HU, IE, IT, LU, MC, NL, PT, RO, SE, SI, SK, TR, BF, BJ, CF, CG, CI, CM, GA, GN, GQ, GW, ML, MR, NE, SN, TD, TG			
CA 2498798	AA	20040401	CA 2003-2498798	20030922
US 2005020613	A1	20050127	US 2003-667570	20030922
EP 1545468	A2	20050629	EP 2003-754790	20030922
R:	AT, BE, CH, DE, DK, ES, FR, GB, GR, IT, LI, LU, NL, SE, MC, PT, IE, SI, LT, LV, FI, RO, MK, CY, AL, TR, BG, CZ, EE, HU, SK			
NO 2005001854	A	20050615	NO 2005-1854	20050415
PRIORITY APPLN. INFO.:			US 2002-412217P	P 20020920
			WO 2003-US29634	W 20030922



## Refine Search

### Search Results -

Terms	Documents
L2 and (zinc adj (sulfate or sulphate))	28

Database:

US Pre-Grant Publication Full-Text Database  
 US Patents Full-Text Database  
 US OCR Full-Text Database  
 EPO Abstracts Database  
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 IBM Technical Disclosure Bulletins

Search:

L5





### Search History

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<u>Set</u> <u>Name</u>	<u>Query</u>	<u>Hit</u> <u>Count</u>	<u>Set</u> <u>Name</u> result set
<i>DB=PGPB,USPT,USOC,EPAB,JPAB,DWPI,TDBD; PLUR=YES; OP=OR</i>			
L5	L2 and (zinc adj (sulfate or sulphate))	28	<u>L5</u>
L4	L3 and (poloxamer or (sorbitan near monoester) or SDS or (sodium adj lauryl adj (sulphate or sulfate)) or (sodium adj dodecyl adj (sulfate or sulphate)) or (glyceryl near monooleate))	50	<u>L4</u>
L3	L2 and (gel near10 polymer)	126	<u>L3</u>
L2	L1 and emetic	1793	<u>L2</u>
L1	analgesic	67565	<u>L1</u>

END OF SEARCH HISTORY